
Vegetation and Foodstuff Monitoring

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Introduction

Because pollutants originally released to the soil, air, or water can be transported to vegetation, the sampling and analysis of native vegetation can provide information about the presence and movement of radionuclides in the environment. Vegetation can contribute a radiation dose to humans directly through ingestion or indirectly through human ingestion of products from animals that have consumed it. DOE guidance states that periodic sampling and analysis of vegetation should be performed to determine if there is measurable long-term buildup of radionuclides in the terrestrial environment (U.S. Department of Energy 1991).

Since 1972, vegetation and foodstuff sampling in the vicinity of LLNL and Site 300 has been part of a continuing LLNL monitoring program designed to measure any changes in environmental levels of radioactivity, to evaluate any increase in radioactivity that might have resulted from LLNL operations, and to calculate potential human doses resulting from direct and indirect ingestion of these products. During 1996, LLNL collected and analyzed samples of native vegetation and wine. Potential human doses from these foodstuffs are calculated using the monitoring data and dose models presented in Appendix B.

Tritium is the nuclide of major interest in the LLNL vegetation and foodstuff monitoring program because LLNL has historically released tritium to the air both accidentally and in the course of routine operations. Tritium is likely to move into the environment as tritiated water and can be assimilated easily into vegetation and foodstuff. It can contribute to human radiation dose burdens if it is inhaled or ingested directly or indirectly. Although other radionuclides are used at LLNL, our assessments show that only tritium could be present in vegetation in detectable concentrations.

Methods

Our methods for monitoring vegetation and wine are presented in the following sections.



Vegetation

LLNL collects vegetation samples, usually annual grasses, quarterly from fixed locations in the Livermore Valley, San Joaquin Valley, and Site 300, and then analyzes them for tritium.

Location maps are provided in **Figures 10-1** and **10-2**. These locations have been selected so samples would represent vegetation from: (1) locations near LLNL that could be affected by LLNL operations, (2) background locations where vegetation was similar to that growing near LLNL but was unlikely to be affected by LLNL operations, and (3) areas of known or suspected LLNL-induced contamination. Sampling location NPER

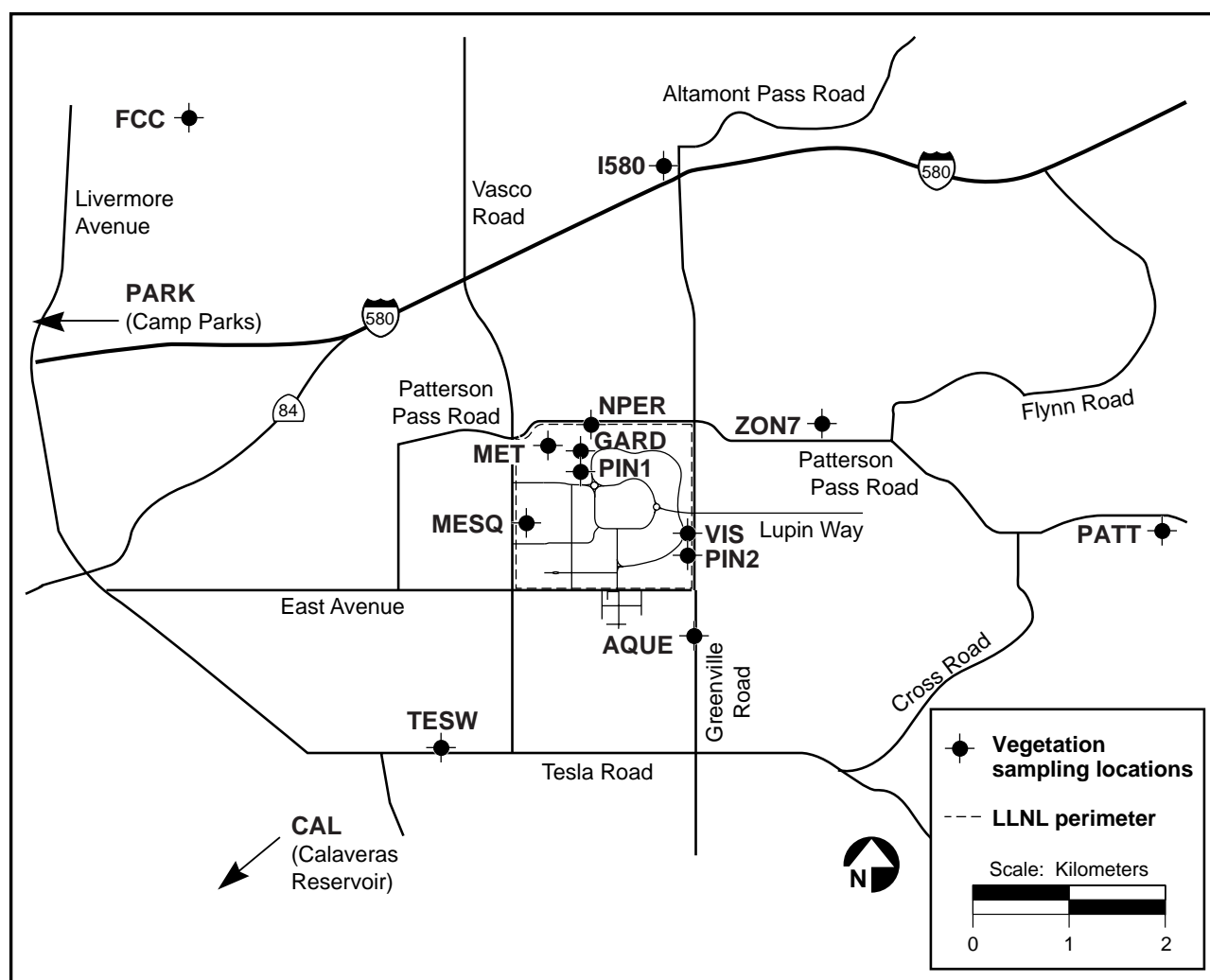


Figure 10-1. Livermore Valley vegetation sampling locations, 1996.

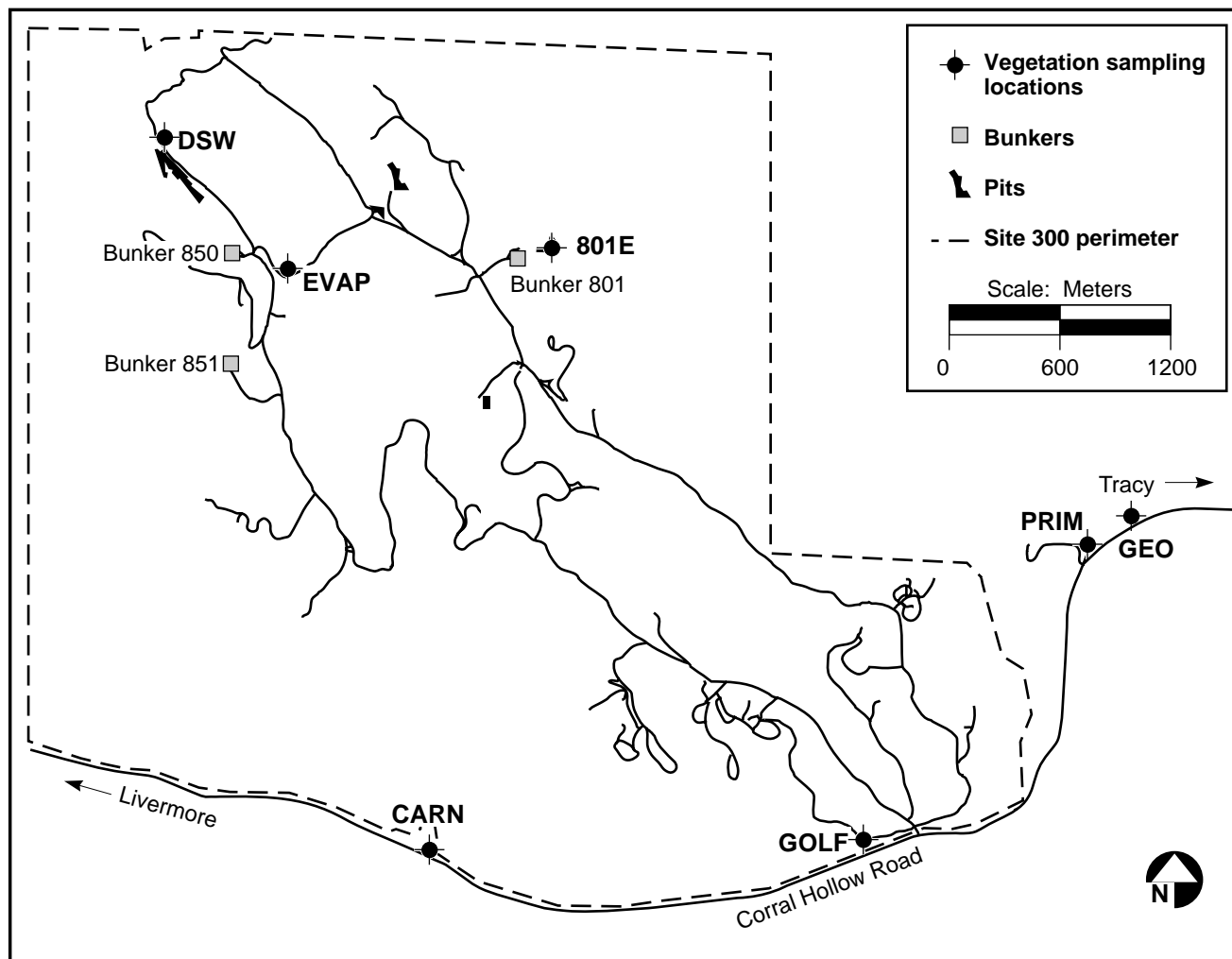


Figure 10-2. Site 300 vegetation sampling locations, 1996.

was added to the sampling network in the first quarter of 1996. It replaced location RAIL, which had become inaccessible. Sampling locations PIN1, PIN2, and PRIM were added in the fourth quarter of 1996. PIN1 and PIN2 were added to evaluate the emissions of tritium from a pine tree that is rooted in tritium-contaminated soil (PIN2 is a tree located in a background location). PRIM is located off site and downwind of Site 300.

All vegetation sampling is conducted according to written and approved standardized procedures (Tate et al. 1995). Approximately 10% of the sites are sampled in duplicate to comply with quality assurance protocols (Garcia and Failor 1993).



Wine

Wine is the most important agricultural product in the Livermore Valley, representing an approximately \$30-million annual industry. Although the tritium concentrations in all wines are low, the data since monitoring began (in 1977) indicate that Livermore Valley wines contain statistically more tritium than do their California counterparts.

Three types of wine samples were collected and analyzed for tritium concentrations: wine produced from grapes grown in the Livermore Valley, wines produced from grapes grown in California outside the Livermore Valley, and wines produced from grapes grown in Europe (France, Germany, and Italy). The latter two groups were divided into 8 and 13 wine-producing regions, respectively, and were used as comparative samples.

The wine samples were purchased from local retailers in a variety of vintages and reflect the body of wines locally available to the general public during 1996. The resulting analytical data can be used to estimate the potential tritium dose received by consumers during the year of purchase. The 1996 sampling data cannot, however, be used to indicate how LLNL's operations affected wines produced in 1996. Some time—in some cases, several years—will have elapsed between the harvest of the grapes and the release of the vintage. However, wine sample data are decay-corrected to original tritium concentrations (given the number of months that have elapsed between wine production and LLNL analysis) to determine trends and to help determine the impact of LLNL operations during a particular vintage year.

The wine samples were submitted for analysis unopened to avoid airborne tritium contamination. Wines were analyzed for tritium using ^3He mass spectrometry in the LLNL Isotope Sciences Noble Gas Mass Spectrometry Laboratory (Surano et al. 1991). This highly sensitive method has a detection limit of less than 0.5 Bq/L (13 pCi/L), and is used to determine the small differences in the tritium content of the samples. Conventional scintillation detection systems typically have detection limits between 5 and 10 Bq/L (150–300 pCi/L); therefore, the differences in the samples would not have been detected had conventional detection methods been used.

Approximately 10% of the total complement of wines was sampled in duplicate, 30% of all the samples were analyzed multiple times, and traceable standards were evaluated to comply with quality assurance protocols.



Results

The results of vegetation and foodstuff monitoring for the Livermore site and Site 300 are presented below.

Livermore site

Vegetation

Table 10-1 shows summary tritium data for vegetation collected in the Livermore site vegetation monitoring program in 1996 (the individual sampling values are presented in Volume 2 of this document). In general, the 1996 tritium levels in vegetation were not significantly different than the levels measured in 1995.

Table 10-1. Tritium in vegetation (in Bq/L), 1996.

Location ^(a)	Detection frequency	Median	Interquartile range	Maximum	Dose ($\mu\text{Sv/y}$) ^(b)	
					Median	Maximum
Livermore site near locations	21/26	5.5	13	96	0.027	0.46
Livermore site intermediate locations	11/16	3.2	3	9.2	0.015	0.044
Livermore site background locations	1/12	<1.9	— ^(c)	2.3	<0.009	0.011
Location DSW at Site 300 ^(d)	3/4	3.6	7.7	32	0.017	0.15
Location EVAP at Site 300 ^(d)	1/4	<1.9	— ^(c)	1360	0.009	6.7
All other locations at LLNL Site 300	1/17	<1.9	— ^(c)	2.0	<0.009	0.009

Note: Detection frequency means the fraction of samples taken having measured values above the detection limit.

^a See **Figures 10-1** and **10-2** for sampling locations.

^b Dose calculated based on conservative assumptions that an adult's diet is exclusively vegetables with this tritium concentration and that meat and milk is derived from livestock fed on grasses with the same concentration of tritium. See Appendix B, Methods of Dose Calculations.

^c Insufficient number of detections to calculate IQR.

^d Sampling location in known area of contamination.

The vegetation locations were put into three groups for statistical evaluation:

- Near—locations at or within 1 km of the Livermore site perimeter. Near locations include AQUE, NPER, GARD, MESQ, MET, PIN1, PIN2, and VIS.
- Intermediate—locations in the Livermore Valley further from the site (1 to 5 km from the Livermore site perimeter) but close enough and often downwind so that they are still potentially under the influence of tritium releases at the site. The intermediate locations were I580, TESW, ZON7, and PATT.



- Far—locations unlikely to be affected by LLNL operations. One background location (CAL) is more than 25 km away. The other two (FCC and PARK) are in the Livermore Valley but are greater than 5 km from the Livermore site and are generally upwind, so they are unlikely to be affected by LLNL operations.

The changes in tritium levels between 1995 and 1996 for the vegetation from within each of the Near, Intermediate, and Far groups were statistically insignificant.

Because the data for tritium in vegetation were lognormally distributed, the means of the logarithms were compared, using the Tukey-Kramer honestly significant difference (HSD) test. This evaluation of the 1996 data showed a significant difference between the Near group and the other two groups; that is, the Near values are significantly different from the Intermediate and Far values, but the Intermediate values are not significantly different from the Far values. **Figure 10-3** shows the historic averages for the three groups. The highest tritium results for individual vegetation sampling locations were found at PIN1, AQUE, and VIS. PIN1 is in a location of known contamination. AQUE and VIS are located downwind of the Livermore site and historically have exhibited higher values than other locations.

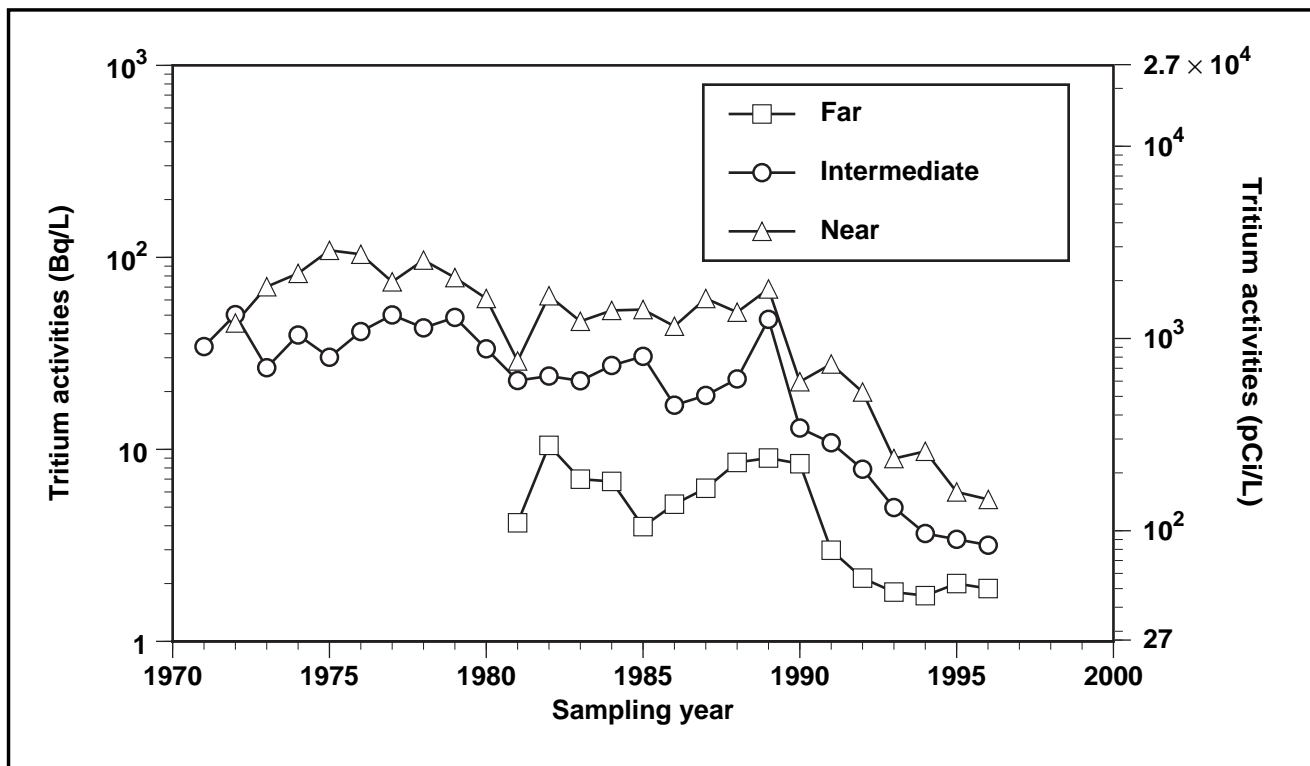


Figure 10-3. Median tritium activities in Livermore Valley vegetation samples, 1971 to 1996.



Wine

The results from the 1996 wine tritium analyses are shown in **Table 10-2**. Tritium concentrations were within the range of those reported in previous years, and they remained low in wines from all areas.

Table 10-2. Tritium (Bq/L) in retail wine, 1996.^(a)

Region	Detection frequency	Median	Interquartile range	Mean	Maximum	Dose ^(b) μSv/y (mrem/y)
Livermore Valley	12/12	3.24	2.56	3.12	5.61	0.0028 (0.00028)
California	6/6	0.58	0.19	0.63	0.93	0.0006 (0.00006)
Europe	4/4	1.52	0.26	1.59	1.92	0.0014 (0.00014)

^a Wines from a variety of vintages were purchased and analyzed during 1996. The concentrations shown are not decay-corrected to vintage year.

^b This dose is calculated from conservative assumption of drinking 52 L wine/year and using the mean concentration of sampled wines.

The data for the 1996 sampling year were analyzed using analysis of variance (ANOVA). The statistical analyses showed that the mean tritium concentration of the Livermore wines sampled was statistically greater than that of the California (other than Livermore) wines. The statistical analyses also indicated that there was no significant difference between the mean tritium values of the European and California wines sampled or between the Livermore and European wines. Multiple comparison tests indicated that the mean levels of the 1996 sampling year data from all areas were not significantly different from those reported for the 1994 and 1995 sampling years. **Figure 10-4**, which shows the results of the wine analyses by sampling year since monitoring began, also shows that 1996 tritium concentrations are among the lowest for all reported Livermore wines.

During the review of historical data in 1995, it was discovered that the data being reported for the 1977 and 1979 sampling years were averages across multiple sampling years. These data have been corrected in **Figure 10-4**, and are the reason for differences observed when comparing this figure to those published before 1995.

Regression analyses and ANOVA of the wine data when decay-corrected and grouped by vintage year (1995 is the last sampled vintage) showed tritium concentrations have statistically decreased for all regions since 1984 (see **Figure 10-5**). Livermore wines, examined by vintage year, had statistically greater tritium concentrations from 1986 through 1994 than both European and California wines. However, the 1995 vintage Livermore wines exhibited slightly lower tritium concentrations than their European counterparts. Whether this is a statistical artifact of small sample size ($n = 2$ for 1995 Livermore wines) or an indication that tritium levels in Livermore wines have actually

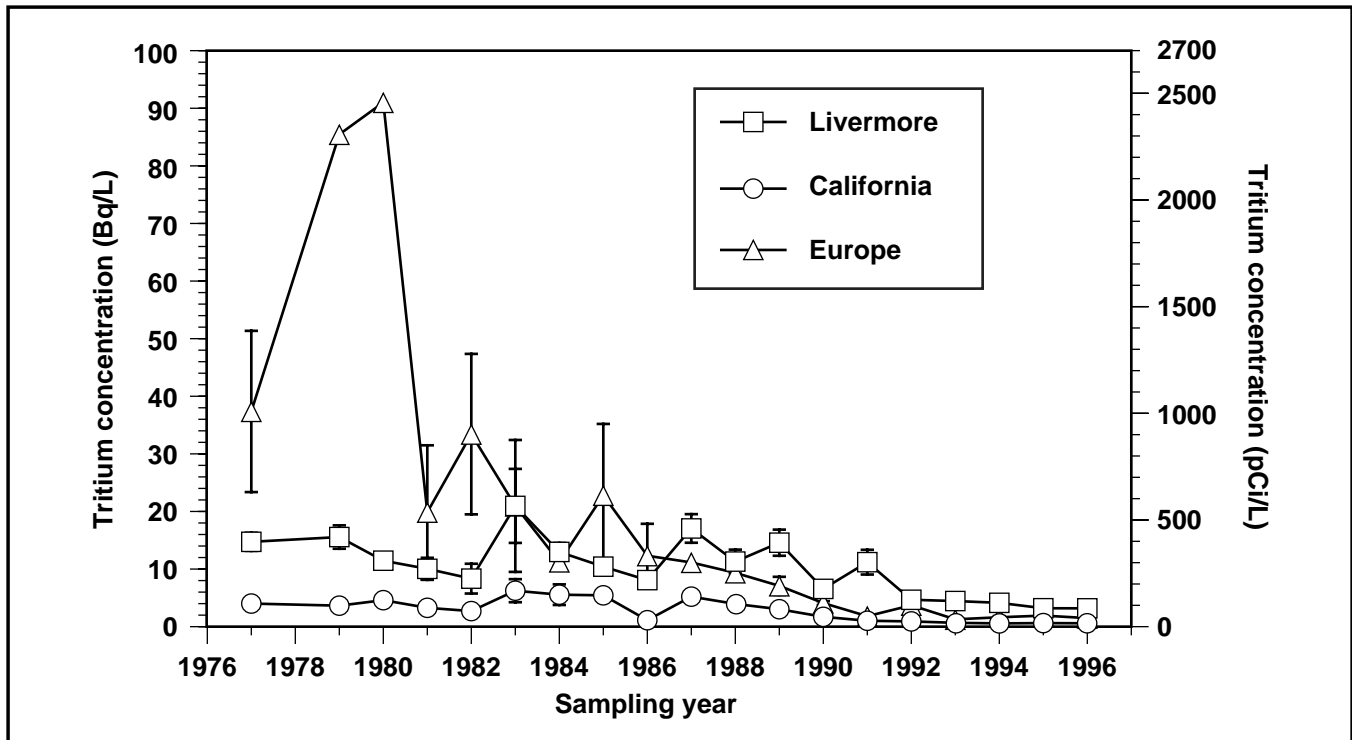


Figure 10-4. Mean tritium in retail wines, 1977 to 1996, plotted by sampling year (error bars are ± 1 standard error).

decreased below that of European wines should be resolved in 1997 when additional 1995 vintage Livermore wines are available for analysis. Nevertheless, it is important to note the continued downward trend in the tritium concentrations of Livermore wines (when decay-corrected and grouped by vintage year) that has been observed since 1984 (when tritium operations at LLNL were scaled down and total amounts of tritium released were reduced).

Site 300

Vegetation

Table 10-1 shows summary tritium data for vegetation collected at Site 300 during 1996. Historic values for tritium at Site 300 sampling locations are shown in Figure 10-6. Of the six sampling locations at Site 300, four yielded results at or near the detection limits. Two locations, EVAP and DSW, yielded results above background.

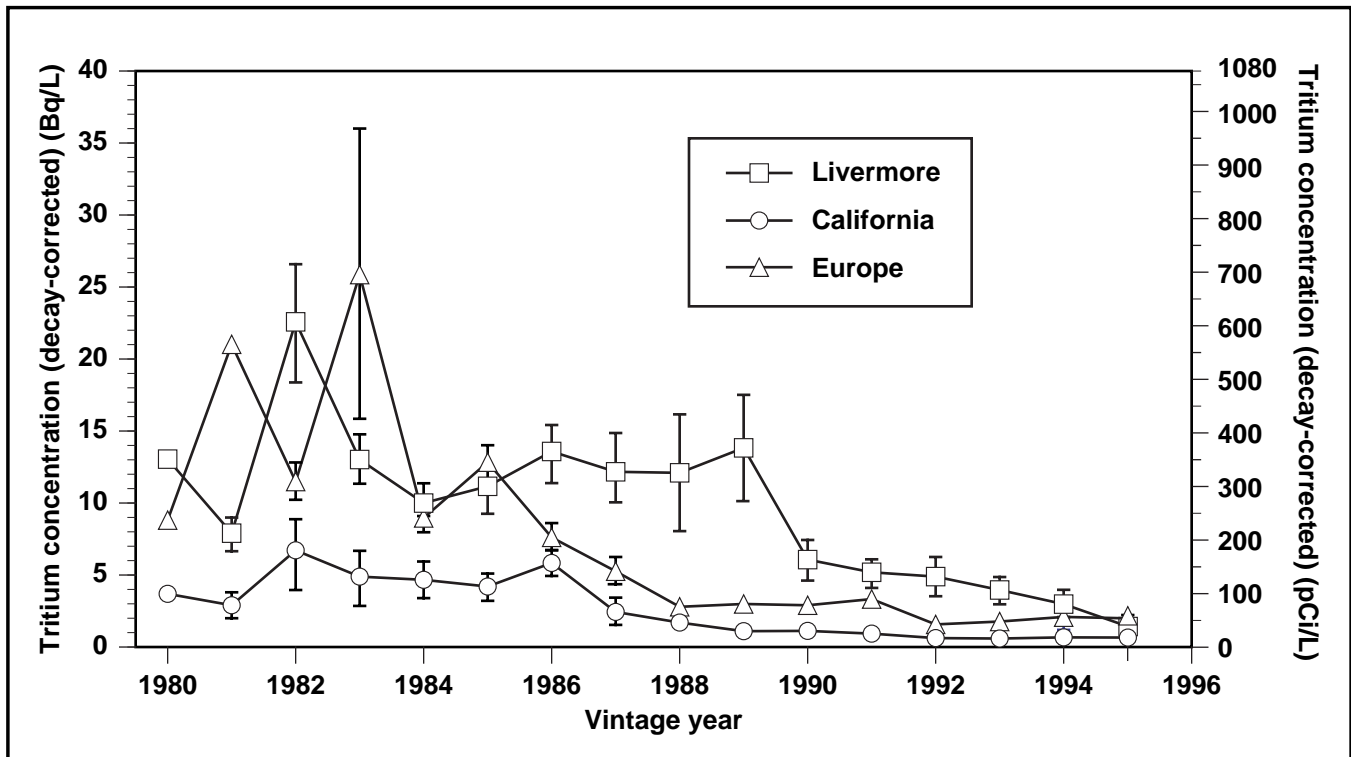


Figure 10-5. Mean tritium in retail wines, vintages 1980 to 1996 values are decay-corrected and plotted by vintage year (error bars are ± 1 standard error).

With one exception, vegetation samples from location DSW contained the highest maximum tritium values detected (see **Table 10-1**). Tritium has been observed in the vegetation of the DSW sampling location since 1971; it is in an area presently being investigated under CERCLA for tritium contamination of ground water. This sampling location is adjacent to a landfill that contains debris contaminated with tritium from past experiments. The landfill area is under continued investigation for tritium in soil and ground water, as described in reports published as part of LLNL's Environmental Restoration Program (Lamarre 1989a, 1989b, and 1989c; Taffet et al. 1989a and 1989b; Taffet et al. 1991; Carlsen 1991a and 1991b; and Webster-Scholten 1994). The highest tritium result for a single vegetation sample occurred at the location EVAP. The sample was not the usual native grass, but a stinging nettle (*Urtica dioica*), which has a relatively long tap root. The location EVAP is near a spring where ground water flows near the surface and evaporates. The ground water in this area is contaminated with tritium which comes from three sources, Pit 3, Pit 5, and the firing table at Building 850 (see discussion of Wells NC7-61 and NC7-69 in Chapter 8, Ground Water). The root of the stinging nettle appears to have reached this ground water, causing the high measured result. Evaluation of the 1996 data using the Tukey-Kramer HSD test on the

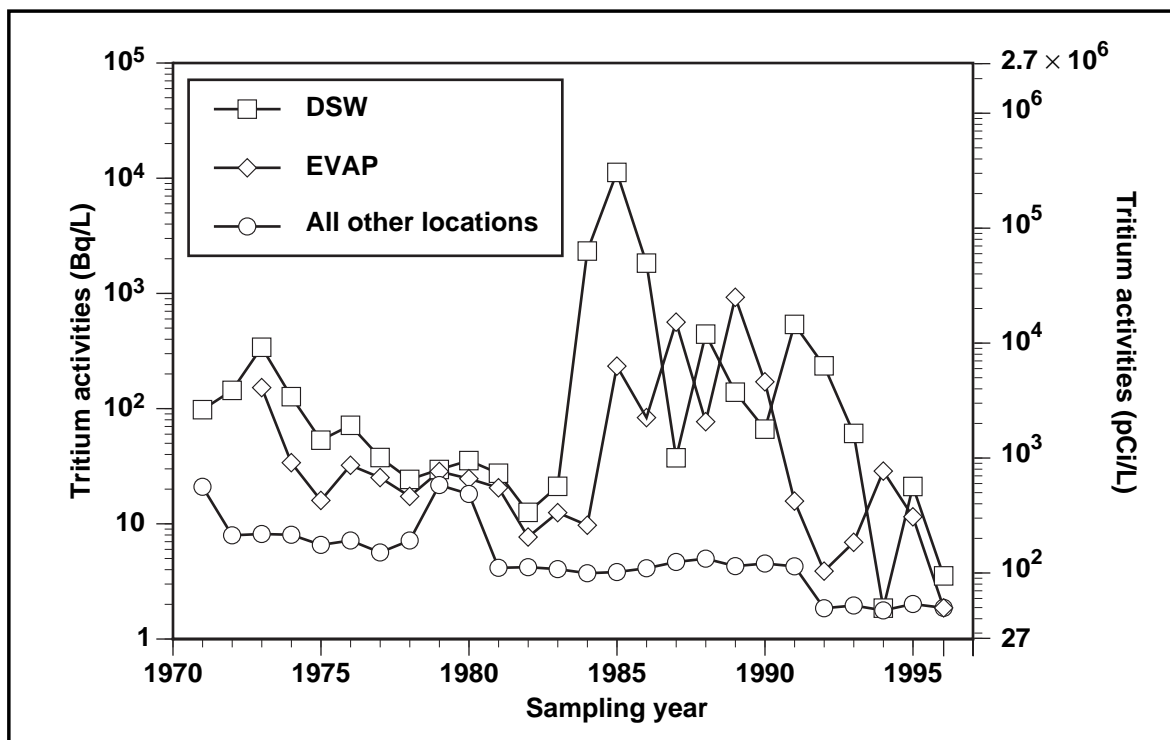


Figure 10-6. Median tritium activities in vegetation at Site 300 sampling locations, 1971 to 1996.

logarithms of the data yielded no significant differences among the various sampling locations; this is a result of the high variability of the data and the low number of data points. However, if the 1995 and the 1996 data are combined, a significant difference is found between the set of locations comprising GEO, CARN, GOLF, and 801E, and locations DSW and EVAP. This is a result of the fact that DSW and EVAP are located in areas of known tritium contamination.

Environmental Impact

The environmental impacts of LLNL operations on vegetation and foodstuff monitoring are small and are presented below for the Livermore site and Site 300.

Livermore Site

LLNL impacts on vegetation in the Livermore Valley remained minimal in 1996. The effective dose equivalents shown in **Table 10-1** were derived using the dose conversion factors provided by DOE (U.S. Department of Energy 1988) and the dose pathway



model from NRC Regulatory Guide 1.109 (U.S. Nuclear Regulatory Commission 1977). Appendix B provides a detailed discussion of dose calculation methods. The dose from tritium in vegetation is based on the conservative assumptions that an adult's diet consists exclusively of vegetables with the measured tritium concentration, and meat and milk derived from livestock fed on grasses with the same concentration. These assumptions are conservative because most vegetables consumed directly by an adult will not contain tritium at the levels reported (the tritium levels will actually be much lower), nor will the livestock actually consume vegetation with the reported levels of tritium. Based on these conservative assumptions, the maximum potential dose (from ingestion of affected vegetation) for 1996 for the Livermore site is $0.46 \mu\text{Sv}$ (0.046 mrem).

No health standards exist for radionuclides in wine. However, all the wine tritium levels were far below drinking water standards. In fact, even the highest detected Livermore Valley value (5.61 Bq/L or 152 pCi/L) represents only 0.8% of the California drinking water standard (740 Bq/L or $20,000 \text{ pCi/L}$). Doses from wine consumption can be calculated according to methods for water ingestion, which are detailed in Appendix B.

The annual dose that corresponds to the highest detected 1996 Livermore Valley tritium value in wine (5.61 Bq/L [152 pCi/L]) is $0.070 \mu\text{Sv}$ (0.0070 mrem), based on the extremely conservative assumption that wine is consumed in the same quantities as water (730 L/year or 2 L/day). Using a more realistic wine consumption factor (52 L/year or 1 L/week of wine from a single area) and the mean tritium values detected in wines from the three sampling areas, the annual dose from Livermore wine would be $0.0028 \mu\text{Sv}$ (0.00028 mrem), from European wine would be $0.0014 \mu\text{Sv}$ (0.00014 mrem), and from California wine would be $0.0006 \mu\text{Sv}$ (0.00006 mrem). Compared with an annual background dose of approximately $3000 \mu\text{Sv}$ (300 mrem), which includes radon, and a $100\text{-}\mu\text{Sv}$ (10-mrem) dose from a typical chest x-ray (Shleien and Terpilak 1984), the potential dose from consuming wine from any area is minute. Therefore, although Livermore wines contained statistically more tritium than wines produced in other areas of California, the effects of the tritium are negligible.

Site 300

In general, LLNL impacts on vegetation at Site 300 for 1996 were insignificant. Tritium levels found in the Site 300 vegetation were comparable to those observed in previous years. With the exception of vegetation from previously identified sites of contamination, the levels were low, near the limits of detection. The areas where tritium is known to be present in the subsurface soil are well delineated and localized.

The calculated maximum potential annual dose from vegetation at sampling location EVAP, based on the maximum value of 1360 Bq/L (38000 pCi/L), is $6.7 \mu\text{Sv}$ (0.67 mrem).



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This dose, which would never actually be received by anyone, is about 15 times less than a chest x-ray (Shleien and Terpilak 1984). This calculation uses the same conservative pathway modeling assumptions, as described above. In actuality, this dose never would be received because vegetation at Site 300 is not consumed by people or by grazing livestock. In comparison, the calculated potential annual dose from vegetation at all other locations at Site 300 had a median value of $<0.009 \mu\text{Sv}$ ($<0.0009 \text{ mrem}$; the value is a “less than” value because all measured tritium levels were less than the detection limit). Tritium levels in vegetation at Site 300 will continue to be monitored.